

**K872838 NORFOLK NORPORT (TM) -SP (SKIN PARALLEL)**Aug 25, 1987  
36 days to decision

K872838 · Product code: LJT · General Hospital

Source: <https://www.510kdatabase.net/k872838/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Jul 20, 1987
Decision date	Aug 25, 1987
Days to decision	36 days
Third-party review	No

**APPLICANT**

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Company	<b>Norfolk Medical Products, Inc.</b>
Location	Walker, MI, US
Contact	MICHAEL J DALTON
510(k) history	20 submissions · 20 cleared · 1983-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k872838/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 18, 2026